## **Subrecipient Statement of Collaborative Intent**

## Part I: To be completed by all subrecipients/subcontractors

All subrecipients, as well as potential subcontractors who anticipate funding under a federal or non-federal contract, should complete this form when participating as a subrecipient or subcontractor in a proposal submitted by the University of Michigan. This form provides a checklist of documents and certifications required by prime sponsors,

and if it is completed, it must be endorsed by the subrecipient's authorized institutional  1. UNIVERSITY OF MICHIGAN PROPOSAL INFORMATION - PASS-TI	
PTE PI:	Prime Sponsor:
PTE Proposal Number:	If Prime Sponsor is NIH: (1) Is this an MPI project? Yes No
	(2) Is the subrecipient a foreign entity? Yes No
2. SUBRECIPIENT PROPOSAL INFORMATION	
Legal Name:	Authorized Official Name:
Address:	Authorized Official Email:
7.444.655.	Authorized Official Phone #:
Phone Number:	Financial Contact Name:
Thore Humber.	Financial Contact Email:
Subrecipient PI:	Project Title:
Subrecipient PI Email:	
Place of Performance:	Project Period:
Congressional District:	Total Request Amount:
UEI Number:	Subrecipient Proposal:
3. PROPOSAL DOCUMENTS	
The following documents are included in our subaward proposal a	nd covered by the certifications below:
Scope of Work (Required)	Facilities & Other Resources
Budget and Justification (Required)	Cost Sharing Amount (if applicable):
Biographical Sketches	Other:
4. CERTIFICATIONS	
Documentation of Subrecipient's approval(s) may be required	
Subrecipient's Scope of Work Includes:	Recombinant DNA
Human Subjects	Dual Use Research of Concern (DURC) For a list of
If human subjects are involved, have all key personnel	applicable agents, see page 9 of NIH policy.
completed human subjects training? Yes No	Large Scale Human or Non-Human Genomic Data (if NIH)
Vertebrate Animals	For applicability, please refer to NOT-OD-14-124. Documentation of an
Custom made antibodies	approved consent form and Institutional Certification will be required prior to the award, at the "Just in Time" stage.
Human induced pluripotent or Human Embryonic Stem Cells	to the dward, at the Bust III Time Stage.
5. FDP Clearinghouse	
Does the subrecipient entity participate in the FDP Clearinghouse	2? Yes No
If <b>YES</b> , complete this page and sign below. (Do not complete Part II of this form.)	
If <b>NO</b> , complete <i>both</i> Parts I and II of this form and sign below.	
If the Prime Awarding Sponsor is the National Institutes of Health, we the	subrecipient organization, agree to abide by all requirements of the NIH Final

Updated Policy Guidance for Subaward/Consortium Written Agreements (NOT-OD-23-182), and will provide access to copies of all lab notebooks, all data, and all documentation that supports the research outcomes as described in the progress report, to the primary recipient with a frequency of no less than once per year, in alignment with the timing requirements for Research Performance Progress Report (RPPR) submission. Such access may be entirely electronic.

By signing below, I certify that I am an authorized institutional representative and the information and representations made herein are true, accurate and complete. The appropriate programmatic and administrative personnel involved in this application are aware of all sponsor policies regarding subawards and are prepared to establish the necessary inter-institutional agreements consistent with those policies. Any work begun and/or expenses incurred prior to execution of a subaward agreement are at the subrecipient's own risk.

Name and Ti	tle of Subrecipient's Authorized Institutional Official
	Regents of the University of Michigan Office o Research and Sponsored Projects Last updated - October 3, 2023
	Name and Ti

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Part II: To be Completed by Subrecipients/Subcontractors NOT participating in the FDP Clearinghouse Pilot See list of participating FDP organizations.

Ceri	ifications
1.	Facilities & Administrative Rates included in this proposal have been calculated based on the following:
Τ.	Our federally negotiated F&A rate for this type of work.
	No federal negotiated rate and we hereby agree to accept the 10% de minimis MTDC rate as a subrecipient.
	In the case of NIH: NIH will continue to reimburse F&A costs to foreign and international organizations at a rate of 8 percent of
	modified total direct costs (MTDC) less only equipment.
	A reduced F&A rate dictated by the prime sponsor that we hereby agree to accept. Rate:
	Not applicable (no indirect cost are requested). If checked, please specify rationale in Comment Section below.
	Indirect costs are not separately requested as costs are fully burdened.
2.	Fringe Benefit Rates included in this proposal have been calculated based on the following:
	Rates are consistent with our federally negotiated rates.
	Other rates (please specify in Comment Section below the basis on which the rate has been calculated) Fringe Benefits are not separately requested as costs are fully burdened.
3.	Financial Conflict of Interest
	Subrecipient organization/institution has an active and enforced policy on conflict of interest consistent with the sponsor's requirements. Examples:
	DOE follows FAL 2022-02
	<ul> <li>NASA follows GCAM Section 3.3</li> <li>NSF follows PAPPG Part II Chapter IX.A</li> </ul>
	PHS follows the provision of 42 CFR Part 50 Subpart F
	Subrecipient does not have an active and/or enforced conflict of interest policy and hereby agrees to abide by U-M's
	policy. To comply with U-M's policy, please attach a completed Non-U-M Affiliate Disclosure Form for each
	investigator on this project.
4.	Ethics in Research Training
	Applicable to projects funded by NSF or any other programs requiring Ethics in Research Training.
	Not applicable because this project is not being funded by NSF or any other programs requiring Ethics in Research Training.
	Subrecipient organization/institution will ensure that all undergraduates, graduate students, and postdoctoral researchers who will be supported by this NSF proposal will be trained on the oversight in the responsible and ethical conduct of research.
5.	Debarment, Suspension, Proposed Debarment
	Are the Subrecipient Institution or any of its principals who are participating in this project debarred, suspended, proposed
	for debarment, declared ineligible or voluntarily excluded from receiving funds from any federal department or agency, or delinquent on any federal debt? Yes No
	If <b>YES</b> , please explain in Comment Section below.
	If <b>NO</b> , the Organization Certifies it (answer all questions below):
	is not presently debarred, suspended, proposed for debarment, or declared ineligible for award of federal contracts;
	is not presently indicted for, or otherwise criminally or civilly charged by a government agency;
	has has not within three (3) years preceding this offer, been convicted of or had a civil judgment rendered against them for commission of fraud or criminal offense in connection with obtaining, attempting to obtain, or performing a public (federal,
	state, or local) contract or subcontract; violation of Federal or State antitrust statutes relating to the submission of offers; or
	commissions of contract or subcontract; violation of Federal or State antitrust statutes relating to the submission of offers; or
	commission of embezzlement, theft, forgery, bribery, falsification, or destruction of records, making false statements or receiving stolen property;
	has has not within 3 years preceding this offer, had one or more contracts terminated for default by any federal agency.

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Audit Status
<ol> <li>Was the subrecipient required to conduct an annual audit in accordance with the Single Audit Act or Uniform Guidance Subpart         <ul> <li>F, Audit Requirements for the most recent Audit year?</li> <li>Yes</li> <li>No</li> </ul> </li> <li>a) Was an audit in accordance with the Single Audit Act completed for the most recent fiscal year?</li> <li>Yes</li> <li>No</li> </ol>
If <b>YES</b> is checked, a complete copy of subrecipient's most recent audit report, or the Internet URL link to a complete copy, must be furnished to U-M before a subaward will be issued. URL:
If <b>no audit was completed OR if Subrecipient is not subject to the Single Audit Act or Uniform Guidance</b> , a limited-scope audit will be reviewed and approved before a subaward can be issued.
b) Were there any audit findings reported? Yes No If <b>YES</b> , please clarify in the comment section below.
Subrecipient Institutional Information Federal policy requires subrecipients of federal funds to be registered in SAM
1. Is subrecipient currently registered in Central Contractor Registration via <a href="SAM">SAM</a> ? Yes No If <b>NO</b> , organizations that have not registered with CCR will need to obtain a Unique Entity Identifier (UEI) number first and then access the CCR online registration through the SAM (System for Award Management) <a href="https://home.page">home.page</a> (U.S. organizations will also need to provide an Employer Identification Number from the Internal Revenue Service that may take an additional 2-5 weeks to become active). Completing and submitting the registration takes approximately one hour to complete and your CCR registration may take 3-5 business days to process. Subrecipient <i>must</i> maintain current CCR information in SAM.
2. EIN #:
3. Be sure that the UEI number is entered on page 1.
Comment